

# Mutagenic potential of *Cordia ecalyculata* alone and in association with *Spirulina maxima* for their evaluation as candidate anti-obesity drugs

R.P. Araldi<sup>1,2</sup>, B.M. Rechiutti<sup>1</sup>, T.B. Mendes<sup>1,3</sup>, E.T. Ito<sup>1</sup> and E.B. Souza<sup>1</sup>

<sup>1</sup>Laboratório de Biologia Molecular, Genética e Mutagenese, Universidade Estadual Paulista “Júlio de Mesquita Filho”, Assis, SP, Brasil

<sup>2</sup>Programa de Pós-Graduação Interunidades em Biotecnologia, Universidade de São Paulo, São Paulo, SP, Brasil

<sup>3</sup>Programa de Pós-Graduação em Biologia Estrutural e Funcional, Universidade Federal de São Paulo, São Paulo, SP, Brasil

Corresponding author: E.B. Souza

E-mail: [edislane@assis.unesp.br](mailto:edislane@assis.unesp.br)

Genet. Mol. Res. 13 (3): 5207-5220 (2014)

Received October 28, 2013

Accepted June 4, 2014

Published July 7, 2014

DOI <http://dx.doi.org/10.4238/2014.July.7.14>

**ABSTRACT.** Obesity is one of the most important nutritional disorders, and can be currently considered as an epidemic. Although there are few weight reduction drugs available on the market, some new drug candidates have been proposed, including *Cordia ecalyculata*, a Brazilian plant with anorectic properties, and *Spirulina maxima*, a cyanobacterium with antioxidant and anti-genotoxic activity. In this study, we evaluated the mutagenic potential of *C. ecalyculata* at doses of 150, 300, and 500 mg/kg alone and in association with *S. maxima* at doses of 75, 150, and 250 mg/kg, respectively, through an *in vivo* micronucleus test, using mice of both sexes, and an *in vitro* micronucleus test and comet assay, using human peripheral blood. For all tests, cyclophosphamide was used as a positive control. The results showed that treatment of 300 mg/kg *C. ecalyculata* and the combination treatment of 500 mg/kg *C. ecalyculata* with 250 mg/kg *S. maxima*

resulted in anorectic effects. The mutagenic tests did not reveal any clastogenic or genotoxic activity for any treatment, indicating that these candidates could be marketed as weight-reduction drugs. Moreover, the drugs contain chemo-preventive substances that can protect against tumorigenesis, which has been associated with obesity.

**Key words:** *Cordia ecalyculata*; *Spirulina maxima*; Micronucleus; Comet assay; Mutagenesis

## INTRODUCTION

With increased availability of high-energy foods in combination with a tendency toward more sedentary lifestyles and genetic factors, obesity has emerged as one of the most important nutritional disorders, and can be currently considered an epidemic (Francischi et al., 2000; Flier and Maratos-Flier, 2013). Obesity is a non-transmissible chronic disease that is characterized by excess body fat, and has multifactorial causes (Scarpato et al., 2011; Flier and Maratos-Flier, 2013).

According to the World Health Organization (WHO), in 2011 there were 1.6 billion adults and 20 million children classified as overweight, and 400 million people were classified as obese worldwide. The WHO also estimated that by 2015, these numbers will likely increase to 2.5 billion overweight and 700 million obese individuals worldwide. In particular, a significant increase in the incidence of obesity has been documented in Brazil in conjunction with a decrease in malnutrition; in 2003, 40% of the Brazilian population was considered to be overweight (Batista-Filho and Rissin, 2003).

This scenario is alarming and constitutes an important public health concern because obesity is associated with metabolic, cardiovascular, respiratory, and hepatobiliary diseases as well as malignancies, including uterine, cervical, breast, ovarian, endometrial, kidney, gastric, esophageal, liver, gallbladder, and pancreatic cancers. Therefore, obesity can generate significant costs to the public health system, contributing to 20% of malignancies in women and 14% in men (Francischi et al., 2000; Flier and Maratos-Flier, 2013; Boeing, 2013; Lehnert et al., 2013; Gilbert and Slingerland, 2013; Williams, 2013).

Although pharmacotherapy is indicated for individuals with a body mass index greater than 27 kg/m<sup>2</sup>, there are few weight-reducing drugs available on the market (Francischi et al., 2000; Kushner, 2013). The drugs most commonly exploited for appetite suppression have two basic pathways of action: 1) central-acting anorectic drugs (sibutramine), which enhance the neurotransmission of the monoamines norepinephrine, serotonin, and dopamine; and 2) classic adrenergic agents such as benzphetamine, phendimetrazine, diethylpropion, mazindol, and phentermine (Kushner, 2013).

However, these substances lead to a diversity of side effects such as increased anxiety, drowsiness, gastrointestinal disorders, and cardiac arrhythmias (Francischi et al., 2000; Kushner, 2013), indicating an urgent need for the identification of new weight-reducing agents. Indeed, it is estimated that the epidemic control of obesity could prevent approximately 97,000 deaths resulting from obesity-associated cancers in the United States alone (Gilbert and Slingerland, 2013). Among the new candidate drugs that could help in weight reduction are extracts of *Cordia ecalyculata* and the cyanobacteria *Spirulina maxima*.

*C. ecalyculata* Vell. (synonym *C. salicifolia* Cham.), popularly known as “porangaba”, “buggy tea”, or “coffee bush”, is a shrub with a height of 8-10 m and a diameter of 30-40 cm belonging to the family Boraginaceae that is distributed throughout the Brazilian territory, and is also found in the rainforests of Argentina and Paraguay (Menghini et al., 2008; da Silva et al., 2010). Its extract is marketed in Brazil and internationally under the name *Pholia magra* and has diuretic activity, leading to appetite suppression and weight reduction (Menghini et al., 2008; da Silva et al., 2010).

Previous studies revealed that *C. ecalyculata* extract shows inhibitory effects against herpes simplex virus type I (Hayashi et al., 1990), and has cardiotonic properties (Matsunaga et al., 1997), hypolipidemic effects (Siqueira et al., 2006), and cytotoxicity against certain types of tumor cells (Arisawa, 1994). The plant is characterized by high concentrations of caffeine, potassium, allantoic acid, and allantoin (Menghini et al., 2008).

*S. maxima* Setchet Gardner is a non heterocystic, filamentous, planktonic, and photosynthetic cyanobacterium (Pandey and Tiwari, 2010) belonging to the order Nostocales, family Oscillatoriaceae, and is characterized by its 65% protein and 30% essential amino acids contents, in addition to containing chelated minerals as iron, potassium, and magnesium, as well as vitamins such as biotin, cyanocobalamin, calcium pantothenate, folic acid, inositol, thiamine, alpha-tocopherol, pyridoxine and riboflavin, phenolic acids, and beta-carotene, and fatty acids such as linoleic and gamalinoleic acid (Dalay, 2002).

*S. maxima* has anti-inflammatory, analgesic, antioxidant, antiviral, cytostatic, lipolytic, hepatoprotective, keratolytic, nutritive, and hypolipidemic activities, and is therefore considered to be one of the greatest discoveries of the 21st century in the field of nutrition (Ponce-Canchihuamán et al., 2010). Furthermore, studies show that the genus *Spirulina* has anti-carcinogenic and anti-genotoxic properties (Miranda et al., 1998; Mishima et al., 1998; Mittal et al., 1999).

The use of herbal medicine aims to prevent, cure, or minimize disease and its symptoms. However, the transformation from a natural product, such as *C. ecalyculata* and *S. maxima*, into a medicinal product should aim at preserving the chemistry and pharmacological integrity while ensuring the maintenance of its biological action and safety for consumption. Thus, toxicological genetic tests are essential to guarantee the safety of drug candidates before they can enter the international market to ensure that they do not confer a risk of carcinogenesis (Witte et al., 2007).

Among the available genetic toxicological tests, the micronucleus test (MNT) and the comet assay are considered to be gold-standard cytogenetic methods (Flores and Yamaguchi, 2008). The MNT was developed by Schmid et al. (1975) and was first used in erythrocytes obtained from the bone marrow of mice (Ribeiro et al., 2003). This test is widely accepted by international agencies and government institutions as part of a battery test for the assessment of new chemicals or drug candidates prior to entering the market (Ribeiro et al., 2003). Therefore, the MNT constitutes a major noninvasive cytogenetic technique that can detect chromosomal mutations, which are the key mechanism leading to carcinogenic events (Fenech, 2000; Carbonari et al., 2005).

However, the experience gained over several decades of toxicology studies reveals that one test is not sufficient to detect all types of mutagenic effects (Witte et al., 2007). Furthermore, a negative result obtained by a test conducted to investigate chromosomal damage in rodent hematopoietic cells is not sufficient to conclude that the component is not mutagenic to somatic cells. Therefore, based on the correlations between results obtained in rodents and humans, the US Food and Drug Administration (FDA) recommended additional testing, both *in vitro* and *in vivo* (Witte et al., 2007).

The comet assay, also known as single cell gel electrophoresis, was introduced by Östling and Johanson in 1984 and later modified by Singh et al. (1988), and possesses the advantages of being a simple and versatile technique that requires only a small number of cells and can be performed with any eukaryotic cell population (Godschalk et al., 2013). In addition, the technique has a wide spectrum of DNA damage detection, and can therefore serve as an integrator test and a complementary assay to other tests (Godschalk et al., 2013).

Thus, in the present study, we aimed to evaluate the anorectic action and the clastogenic and genotoxic potential of *C. ecalyculata* at three commonly marketed doses (150, 300, and 500 mg/kg). We further evaluated these effects when *C. ecalyculata* was combined with *S. maxima* at respective doses of 75 mg/kg (Ce 150 + Sm 75), 150 mg/kg (Ce 300 + Sm 150), and 250 mg/kg (Ce 500 + Sm 250). An *in vivo* MNT was conducted in polychromatic erythrocytes of mice and two *in vitro* tests were conducted (an MNT and a comet assay) in human binucleated lymphocytes.

## MATERIAL AND METHODS

### *In vivo* MNT in polychromatic erythrocytes and anorectic evaluation

We used a total of 80 mice (*Mus musculus*, albino, Swiss strain), 40 males and 40 females aged between 7 and 11 weeks, obtained from the central biotery of Universidade Estadual Paulista “Júlio de Mesquita Filho” (UNESP). The protocols used were approved by the Ethics Committee on Animal Use of the same university (protocol No. 001194/10). The choice of mice was based on the fact that oncogenetic and functional studies have shown that they do not lose or sequester micronucleated erythrocytes, such as occurs in rats (Ribeiro et al., 2003; Griffiths et al., 2005).

The mice were acclimated in the biotery of UNESP - Faculdade de Ciências e Letras de Assis - for five days under the following conditions: 12:12-h light/dark cycle, 150 lux lighting, temperature of  $22^{\circ} \pm 2^{\circ}\text{C}$ , water and food *ad libitum*. The mice were divided into eight groups of five animals each, separated by sex (Table 1). The analysis of anorectic action was conducted by measuring body weight before and after mice received treatments.

**Table 1.** Different treatments employed, showing doses in mg/kg to *in vivo* micronucleus test (MNT) and  $\mu\text{L}/\text{mL}$  to *in vitro* MNT and comet assay.

Group	Treatment	<i>In vivo</i> MNT Dose (mg/kg)	<i>In vitro</i> MNT Dose ( $\mu\text{L}/\text{mL}$ )	Comet assay Dose ( $\mu\text{L}/\text{mL}$ )
C+	Cyclophosphamide	50	50	50
C-	Saline	-	Not applied	Not applied
Ce 150	<i>C. ecalyculata</i>	150	150	150
Ce 300	<i>C. ecalyculata</i>	300	300	300
Ce 500	<i>C. ecalyculata</i>	500	500	500
Ce 150 + Sm 75	<i>C. ecalyculata</i> + <i>S. maxima</i>	150 + 75	Not tested	150 + 75
Ce 300 + Sm 150	<i>C. ecalyculata</i> + <i>S. maxima</i>	300 + 150	Not tested	300 + 150
Ce 500 + Sm 250	<i>C. ecalyculata</i> + <i>S. maxima</i>	500 + 250	Not tested	500 + 250

The positive control groups (C+) of both sexes were treated with 50 mg/kg cyclophosphamide (Sigma-Aldrich; St. Louis, MO, USA) by intraperitoneal injection, administered in a single dose. The negative control groups (C-) of both sexes were treated with 0.3 mL saline by

gavage, performed twice a day for seven consecutive days. The experimental groups received a final volume of 0.3 mL respective treatment solution (Table 1) through gavage, performed twice a day for seven consecutive days. The mice were sacrificed by cervical dislocation 48 h after the final dose administration. The femurs were removed and the bone marrow was extracted by injecting 1 mL fetal bovine serum (Cultilab; Campinas, SP, Brazil). The material was homogenized on a Petri dish and centrifuged for 5 min at 36.54 g. The pellet was collected and smeared onto 26 x 76-mm slides. The material was fixed in absolute methanol for 5 min and subjected to the following staining protocol: 10 min in 100% Giemsa (eosin methylene blue), washed in distilled water, placed in a solution of 1:5 Giemsa:phosphate buffer, pH 6.0, for 8 min, and washed in xylene. The slides were mounted by gluing a coverslip with Entellan. The material was analyzed under a binocular optical microscope at a total magnification of 1000X. A total of 2000 polychromatic erythrocytes were analyzed per animal, and the frequency of micronucleated polychromatic erythrocytes (MNPCEs) per animal was calculated.

The statistical analyses were performed using the Biostat software version 5.0. The Student *t*-test and the nonparametric Wilcoxon test were used to verify the effectiveness of the drug on weight loss (i.e., change in weight before and after the different treatments), and analysis of variance (ANOVA) was used for comparisons among the treatment groups. The evaluation of the frequency of MNPCEs was performed using the chi-square test, as recommended by Ribeiro et al. (2003). All statistical tests were interpreted at a significance level of 5%.

### ***In vitro* MNT in binucleated lymphocytes**

Five milliliters peripheral blood was collected from 10 volunteers (5 men and 5 women), using a heparinized vacutainer system. The subjects were aged 18-25 years and did not use drugs or any kind of chemical substance, including alcohol and tobacco. The material collected was used in the *in vitro* MNT and comet assay. The protocols were approved by the Ethics Committee in Research of UNESP - Faculdade de Ciências e Letras de Assis (process No. 1679/2010).

The material was immediately processed in order to avoid possible damage to the DNA. For each individual, five cultures were mounted and distributed into five treatment groups, as shown in Table 1. The combination of *C. ecalyculata* and *S. maxima* at three different doses was performed as a complementary test to evaluate the clastogenic action of the extract of *C. ecalyculata* since previous studies demonstrated the anti-clastogenic action of *S. maxima* (Miranda et al., 1998; Mishima et al., 1998; Mittal et al., 1999). Thus, the combination was not tested in the *in vitro* MNT but rather through the comet assay, which has high DNA damage detection sensitivity.

The cultures were performed by addition of 0.2 mL peripheral blood in a culture flask containing 5 mL RPMI 1640 medium, 15% fetal bovine serum, 0.1 mL L-glutamine, and 0.1 mL phytohemagglutinin. The material was incubated at 37°C for 8 h, and then added to 0.1 mL *C. ecalyculata* solutions, respectively (Table 1). After 44 h, 0.2 mL cytochalasin B was added to block cytokinesis. After 72 h, the culture was stopped with the addition of 0.5 mL methanol:acetic acid fixative (3:1) for 5 min at room temperature. The material was centrifuged at 10.39 g and the supernatant was discarded. The pellet was homogenized with 5 mL fixative and centrifuged at 10.39 g. The pellet was aspirated and transferred to slides, which were stained with a 1:3 Giemsa:phosphate buffer solution, pH 6.8, for 8 min.



After staining, coverslips were placed on slides with Entellan® and the material was analyzed in a blind test under a binocular optical microscope to observe the frequency of micronucleated binucleated lymphocytes in a total of 1000 analyzed cells. Statistical analysis was performed using the Kruskal-Wallis test followed by a *post-hoc* Dunn test, both at a significance level of 5%.

### Comet assay

Slides for the comet assay were prepared as follows: slides of 26 x 76 mm were embedded in a solution of normal melting point agarose (NMA; Invitrogen; Carlsbad, CA, USA) diluted in phosphate-buffered solution (PBS) to 1.5% at 60°C, and one side of each slide was wiped clean with a paper towel. The slides were dried in a horizontal position overnight.

The blood samples were collected and fractionated into aliquots of 200 µL each in 0.2-mL polypropylene tubes containing 200 µL solution composed of RPMI 1640 medium with different concentrations of the drugs tested (Table 1). One blood aliquot from each donor, not treated with any compound was used as a negative control, and one aliquot treated with 50 µL/mL cyclophosphamide was used as a positive control. Aliquots were incubated at 37°C for 1 h under constant agitation.

After incubation, the aliquots were centrifuged for 1 min at 504 g, and the supernatant was discarded. Ten microliters obtained pellet was added to 75 µL low melting point agarose, and a final volume of 85 µL was immediately transferred to NMA pre-coated slides. The slides were covered with coverslips and maintained at 4°C for 20 min. The coverslips were gently removed and the slides were placed in a coplin jar containing lysis solution [2.5 mM NaCl, 100 mM ethylenediaminetetraacetic acid (EDTA), 10 mM Tris-HCl, 1.1% Triton X-100, and 11.2% dimethyl sulfoxide] at 4°C for 1 h. Subsequently, all procedures were conducted under dark conditions to prevent the induction of DNA damage.

After lysis, the slides were washed with PBS and transferred to an electrophoresis tank containing electrophoresis buffer (300 mM NaOH and 1 mM EDTA, pH >13) at 4°C for 40 min to induce unwinding of double-stranded DNA. Next, the electrophoretic run was performed with a current of 25 V (0.86 V/cm), 300 mA for 20 min to promote the migration of free DNA fragments toward the anode. The slides were transferred to a coplin jar containing neutralizing buffer (400 mM Tris-HCl, pH 7.5) for 5 min. The material was fixed in absolute ethanol for 5 min.

The slides were stained with 75 µL 20 µg/mL ethidium bromide solution, and analyzed under an epifluorescence microscope (Carl Zeiss Axio Scope A1) equipped with an excitation filter of 510-560 nm and barrier of 590 nm, at a total magnification of 400X. One hundred nucleoids were analyzed per slide, which were scored on a scale of 0 (without DNA damage) to 3 (maximum DNA damage), according to Tug et al. (2011). The scores were obtained by summing the product of the observed number of nucleoids per class by its respective class value. Statistical analysis was performed using the Kruskal-Wallis test followed by the *post-hoc* Dunn test, both with a significance level of 5%.

## RESULTS

### Analysis of the anorectic action

The mice were weighed before and after the treatments and the results are shown in

Table 2. The Shapiro-Wilk test revealed that the samples were normally distributed ( $P > 0.05$ ), except for the combination treatment group Ce 500 + Sm 250 of females ( $P = 0.0183$  and  $P = 0.0464$  before and after treatment, respectively). Based on these data, we performed the Student *t*-test for paired samples that were normally distributed and the nonparametric Wilcoxon test for the group of females that did not show normality, both at a significance level of 5%. The results indicated significant differences in weight for groups C- and Ce 500 + Sm 250 of males and C+, C-, Ce 150, Ce 500, Ce 150 + Sm 75, and Ce 300 + Sm 150 of females (Table 3). In both sexes, there was an increase in weight for the negative control groups (C-), as expected. An increase in weight was observed in the positive control group (C+) of female mice.

**Table 2.** Body weight before and after the different treatments, of ten mice, being five male and five female, followed by the values of mean and stand deviation (SD).

	C+		C-		Ce 150		Ce 300		Ce 500		Ce 150 + Sm 75		Ce 300 + Sm 250		Ce 500 + Sm 250	
Male	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After
1	24.2	23.7	28.8	32.1	27.2	25.1	27.8	27.6	29.0	29.5	34.5	35.9	40.2	39.4	39.9	38.9
2	27.7	27.8	32.7	38.6	27.6	28.2	31.0	35.3	33.1	32.9	35.7	34.7	43.8	43.8	43.0	42.9
3	29.0	30.5	35.7	44.6	30.3	33.1	36.2	35.8	36.9	33.3	35.8	37.7	44.3	46.0	47.1	46.0
4	29.8	32.4	37.5	44.8	33.0	38.8	36.2	36.6	40.9	42.7	37.3	37.7	48.0	49.9	47.7	46.4
5	30.1	35.6	38.2	45.1	36.8	39.9	33.1	37.8	44.0	46.0	40.3	40.7	48.9	52.9	55.3	54.7
Means	28.1	30.0	34.5	41.0	30.9	33.0	32.8	34.6	36.7	36.8	36.7	37.3	45.0	46.4	46.6	45.7
SD	2.4	4.5	3.8	5.6	4.0	6.4	3.5	4.0	5.9	7.0	2.2	2.2	3.5	5.2	5.8	5.8
Female	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After	Before	After
1	21.8	22.7	23.3	26.8	26.7	21.7	23.2	22.7	31.7	29.7	31.8	28.0	36.2	35.2	35.7	35.1
2	24.3	24.6	23.4	27.4	28.6	24.9	26.8	26.8	34.3	30.2	35.5	31.9	37.9	37.8	38.4	36.0
3	27.9	34.6	25.7	30.5	29.0	25.2	29.5	28.7	36.7	33.6	38.6	37.3	40.6	39.3	39.2	39.6
4	31.8	38.3	25.8	30.8	29.3	26.8	31.1	31.4	37.5	35.2	38.8	39.0	41.9	41.2	39.4	40.0
5	36.3	40.5	30.9	31.1	32.1	27.8	34.3	32.9	40.1	39.1	40.3	39.7	42.5	41.6	56.9	57.3
Means	28.4	32.1	25.8	29.3	29.1	25.2	28.9	28.5	36.0	33.5	37.0	35.1	39.8	39.0	41.9	41.6
SD	5.8	8.0	3.0	2.0	1.9	2.3	4.2	4.0	3.1	3.8	3.3	5.0	2.6	2.6	8.5	9.0

**Table 3.** Statistic results of weight reduction before and after the different treatments for male and female mice, followed by the P valor.

Treatments	Male		Female	
	Statistic values	P	Statistic values	P
C+	-1.7312	0.0792	-2.7500	0.0256
C-	-6.9757	0.0011	-4.0307	0.0079
Ce 150	-1.5410	0.0991	9.4007	0.0004
Ce 300	-1.5602	0.0968	1.6054	0.0918
Ce 500	-0.0989	0.4630	-4.7847	0.0044
Ce 150 + Sm 75	-1.2440	0.1407	2.2637	0.0431
Ce 300 + Sm 150	-1.6318	0.0890	4.0000	0.0081
Ce 500 + Sm 250	3.8485	0.0092	t/Wilcoxon = 26.0000	0.3770

The experimental groups showed a weight reduction in males treated with the maximum combination dose (Ce 500 + Sm 250) and in females treated with *C. ecalyculata* alone at doses of 150 and 500 mg/kg and in combination with *S. maxima* at doses of 150 + 75 and 300 + 150 mg/kg. In order to compare the anorectic action of different treatments, ANOVA was performed followed by the *post-hoc* Tukey test, revealing significant differences in weight between groups among the male ( $F = 4.8723$ ,  $P = 0.001$ ) and female ( $F = 14.0667$ ,  $P = 0.000$ ) mice.

The anorectic effect was similar among the Ce 300, Ce 500, Ce 150 + Sm 75, Ce 300 + Sm 150, and Ce 500 + Sm 250 groups in males, and between the Ce 300 and Ce 500 + Sm 250 groups in females (Table 4).

**Table 4.** Weight difference among the different treatments in both male and female mice, in which negative values indicate weight reduction.

Male	C+	C-	Ce 150	Ce 300	Ce 500	Ce 150 + Sm 75	Ce 300 + Sm 150	Ce 500 + Sm 250
1	-0.5 <sup>a</sup>	+3.3 <sup>a</sup>	-2.1 <sup>a</sup>	-0.2 <sup>ab</sup>	+0.5 <sup>ab</sup>	+1.4 <sup>ab</sup>	-0.8 <sup>ab</sup>	-1.0 <sup>ab</sup>
2	+0.1 <sup>a</sup>	+5.9 <sup>a</sup>	+0.6 <sup>a</sup>	+4.3 <sup>ab</sup>	-0.2 <sup>ab</sup>	-1.0 <sup>ab</sup>	0.0 <sup>ab</sup>	-0.1 <sup>ab</sup>
3	+1.5 <sup>a</sup>	+8.9 <sup>a</sup>	+2.8 <sup>a</sup>	-0.4 <sup>ab</sup>	-3.6 <sup>ab</sup>	+1.9 <sup>ab</sup>	+1.7 <sup>ab</sup>	-1.1 <sup>ab</sup>
4	+2.6 <sup>a</sup>	+7.3 <sup>a</sup>	+5.8 <sup>a</sup>	+0.4 <sup>ab</sup>	+1.8 <sup>ab</sup>	+0.4 <sup>ab</sup>	+1.9 <sup>ab</sup>	-1.3 <sup>ab</sup>
5	+5.5 <sup>a</sup>	+6.9 <sup>a</sup>	+3.7 <sup>a</sup>	+4.7 <sup>ab</sup>	+2.0 <sup>ab</sup>	+0.4 <sup>ab</sup>	+4.0 <sup>ab</sup>	-0.6 <sup>ab</sup>
Female	C+	C-	Ce 150	Ce 300	Ce 500	Ce 150 + Sm 75	Ce 300 + Sm 150	Ce 500 + Sm 250
1	+0.9 <sup>a</sup>	+3.5 <sup>a</sup>	-5.0 <sup>b</sup>	-0.5 <sup>bc</sup>	-2.0 <sup>b</sup>	-3.8 <sup>b</sup>	-1.0 <sup>b</sup>	-0.6 <sup>bc</sup>
2	+0.3 <sup>a</sup>	+4.0 <sup>a</sup>	-3.7 <sup>b</sup>	0.0 <sup>bc</sup>	-4.2 <sup>b</sup>	-3.6 <sup>b</sup>	-0.1 <sup>b</sup>	-2.4 <sup>bc</sup>
3	+6.7 <sup>a</sup>	+4.8 <sup>a</sup>	-3.8 <sup>b</sup>	-0.8 <sup>bc</sup>	-3.1 <sup>b</sup>	-1.3 <sup>b</sup>	-1.3 <sup>b</sup>	+0.4 <sup>bc</sup>
4	+6.5 <sup>a</sup>	+5.0 <sup>a</sup>	-2.5 <sup>b</sup>	+0.3 <sup>bc</sup>	-2.3 <sup>b</sup>	+0.2 <sup>b</sup>	-0.7 <sup>b</sup>	+0.6 <sup>bc</sup>
5	+4.2 <sup>a</sup>	+0.2 <sup>a</sup>	-4.3 <sup>b</sup>	-1.4 <sup>bc</sup>	-1.0 <sup>b</sup>	-0.6 <sup>b</sup>	-0.9 <sup>b</sup>	+0.4 <sup>bc</sup>

Numbers followed by at least one equal letter indicate absence of significant statistic differences.

### *In vivo and in vitro* MNT

The chi-squared test results showed that the number of micronucleated cells *in vivo* significantly increased in the positive control groups of both sexes, as expected. There was no significant difference in the number of micronucleated cells between the negative control and experimental groups, thus confirming that the drugs, at the tested doses, did not show any aneuploidogenic and/or clastogenic activity *in vivo* (Table 5).

**Table 5.** Number of micronucleated polychromatic erythrocytes observed per animal of both sexes, followed by total.

Treatments	Groups of male mice						E	$\chi^2$	P
	1	2	3	4	5	Total			
C+	7	6	17	7	3	40	8.0	14.000	0.007
C-	4	3	3	0	1	11	2.2	4.909	0.296
Ce 150	3	1	2	3	3	12	2.4	1.333	0.855
Ce 300	3	3	3	3	1	13	2.6	1.269	0.870
Ce 500	0	1	1	2	0	4	0.8	3.500	0.477
Ce 150 + Sm 75	0	2	0	2	3	7	1.4	5.142	0.272
Ce 300 + Sm 150	1	3	4	0	0	8	1.6	8.250	0.082
Ce 500 + Sm 250	0	3	2	1	1	7	1.4	3.714	0.446
Treatments	Groups of female mice						E	$\chi^2$	P
	1	2	3	4	5	Total			
C+	21	6	13	12	8	60	12.0	11.166	0.024
C-	0	0	2	1	0	3	0.6	5.333	0.254
Ce 150	3	3	3	3	3	14	2.8	0.285	0.990
Ce 300	2	4	1	0	2	9	1.8	4.888	0.290
Ce 500	1	0	0	2	2	5	1.0	4.000	0.406
Ce 150 + Sm 75	0	2	2	1	2	7	1.4	2.285	0.683
Ce 300 + Sm 150	4	1	2	2	4	13	2.6	2.769	0.597
Ce 500 + Sm 250	0	0	5	2	2	9	1.8	9.330	0.053

No. of micronucleated cells analysed and expected value (E), Chi-square ( $\chi^2$ ) and probability (P).



The Kruskal-Wallis test, followed by the *post-hoc* Dunn test, revealed that the frequency of micronucleated cells *in vitro* (among 1000 binucleated lymphocytes) differed significantly between the positive and negative control groups, although no differences were observed between the experimental groups and the negative control ( $H = 25.9712$ ,  $P = 0.0000$ ), confirming the *in vivo* MNT results (Table 6). Together, these results demonstrate that *C. ecalyculata* does not have clastogenic activity.

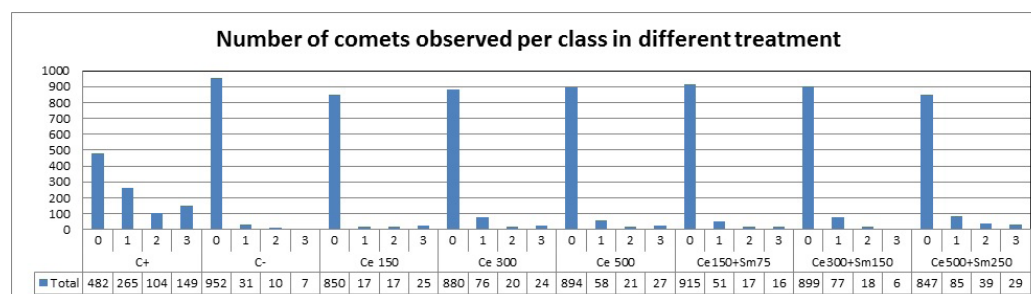
**Table 6.** Number of micronucleated cells observed per treatment, followed by the statistic comparison through Dunn test.

Sample	C+	C-	Ce 150	Ce 300	Ce 500
1	17 <sup>a</sup>	8 <sup>b</sup>	11 <sup>b</sup>	8 <sup>b</sup>	7 <sup>b</sup>
2	16 <sup>a</sup>	7 <sup>b</sup>	1 <sup>b</sup>	1 <sup>b</sup>	1 <sup>b</sup>
3	11 <sup>a</sup>	4 <sup>b</sup>	5 <sup>b</sup>	1 <sup>b</sup>	0 <sup>b</sup>
4	24 <sup>a</sup>	9 <sup>b</sup>	1 <sup>b</sup>	6 <sup>b</sup>	1 <sup>b</sup>
5	12 <sup>a</sup>	3 <sup>b</sup>	2 <sup>b</sup>	0 <sup>b</sup>	3 <sup>b</sup>
6	12 <sup>a</sup>	1 <sup>b</sup>	3 <sup>b</sup>	2 <sup>b</sup>	0 <sup>b</sup>
7	13 <sup>a</sup>	2 <sup>b</sup>	1 <sup>b</sup>	1 <sup>b</sup>	0 <sup>b</sup>
8	21 <sup>a</sup>	4 <sup>b</sup>	5 <sup>b</sup>	17 <sup>b</sup>	2 <sup>b</sup>
9	18 <sup>a</sup>	3 <sup>b</sup>	6 <sup>b</sup>	11 <sup>b</sup>	2 <sup>b</sup>
10	17 <sup>a</sup>	1 <sup>b</sup>	2 <sup>b</sup>	4 <sup>b</sup>	1 <sup>b</sup>
Total	161	42	37	51	17

Number followed by at least one equal letter indicate absence of significative statistic differences.

## Comet assay

A total of 100 nucleoids per treatment were analyzed and given a score of 0 to 3, according to Tug et al. (2011), and the comet score values are shown in Figure 1. The results of the Kruskal-Wallis test, followed by the *post-hoc* Dunn test, revealed that scores differed significantly between the positive and negative control groups, but not among experimental groups, and between experimental groups and the negative control ( $H = 32.7984$ ;  $P = 0.0000$ ) (Table 7). These results confirm the data obtained through the *in vivo* and *in vitro* MNTs, and further validate that the drugs, in the tested doses and in their respective associations, are safe from a mutagenic perspective while still demonstrating anorectic properties. Together, the results show that these drugs are potential candidates to enter into the global market for obesity treatment.

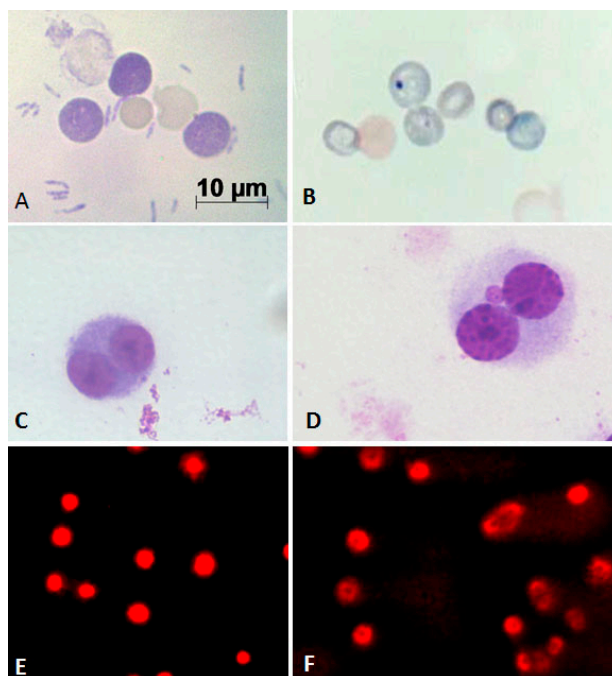


**Figure 1.** Histogram showing the total number of comets observed per class in different treatments.

**Table 7.** Comet score observed per treatment and statistic compasison through Dunn test.

Sample	C+	C-	Ce 150	Ce 300	Ce 500	Ce 150 + Sm 75	Ce 300 + Sm 150	Ce 500 + Sm 250
1	99 <sup>a</sup>	5 <sup>b</sup>	14 <sup>b</sup>	12 <sup>b</sup>	51 <sup>b</sup>	20 <sup>b</sup>	15 <sup>b</sup>	54 <sup>b</sup>
2	52 <sup>a</sup>	11 <sup>b</sup>	25 <sup>b</sup>	15 <sup>b</sup>	49 <sup>b</sup>	6 <sup>b</sup>	18 <sup>b</sup>	36 <sup>b</sup>
3	91 <sup>a</sup>	4 <sup>b</sup>	4 <sup>b</sup>	23 <sup>b</sup>	5 <sup>b</sup>	5 <sup>b</sup>	6 <sup>b</sup>	29 <sup>b</sup>
4	99 <sup>a</sup>	6 <sup>b</sup>	18 <sup>b</sup>	31 <sup>b</sup>	21 <sup>b</sup>	3 <sup>b</sup>	5 <sup>b</sup>	4 <sup>b</sup>
5	82 <sup>a</sup>	7 <sup>b</sup>	3 <sup>b</sup>	25 <sup>b</sup>	10 <sup>b</sup>	3 <sup>b</sup>	5 <sup>b</sup>	10 <sup>b</sup>
6	43 <sup>a</sup>	25 <sup>b</sup>	97 <sup>b</sup>	60 <sup>b</sup>	8 <sup>b</sup>	33 <sup>b</sup>	30 <sup>b</sup>	18 <sup>b</sup>
7	137 <sup>a</sup>	4 <sup>b</sup>	11 <sup>b</sup>	9 <sup>b</sup>	6 <sup>b</sup>	6 <sup>b</sup>	6 <sup>b</sup>	27 <sup>b</sup>
8	103 <sup>a</sup>	6 <sup>b</sup>	20 <sup>b</sup>	3 <sup>b</sup>	3 <sup>b</sup>	5 <sup>b</sup>	14 <sup>b</sup>	31 <sup>b</sup>
9	104 <sup>a</sup>	4 <sup>b</sup>	21 <sup>b</sup>	6 <sup>b</sup>	7 <sup>b</sup>	37 <sup>b</sup>	20 <sup>b</sup>	6 <sup>b</sup>
10	110 <sup>a</sup>	3 <sup>b</sup>	25 <sup>b</sup>	5 <sup>b</sup>	20 <sup>b</sup>	15 <sup>b</sup>	12 <sup>b</sup>	32 <sup>b</sup>

Figure 2 shows images of micronucleated cells and class 3 comets from the respective positive controls and normal cells (i.e., without DNA damage), observed in the experimental groups.



**Figure 2.** Images of normal polychromatic erythrocytes (A) and micronucleated cell (B), normal binucleated lymphocytes (C) and micronucleated cell (D) and comets class 0 (E) and 3 (F).

## DISCUSSION

The increased prevalence of obesity has led to a reduction in the quality of life for a large proportion of the global population, with clear impacts on public health costs; indeed, over 20% of the USA healthcare costs is spent on obesity-associated complications and deaths.

Therefore, the search for new treatments that could assist in weight reduction while not conferring additional health risks is necessary (Scarpato et al., 2011; Boeing, 2013; Lehnert et al., 2013; Gilbert and Slingerland, 2013).

In this regard, the evaluation of DNA damage is an essential part of the characterization of new drug candidates before entrance into the global market (Frieauff et al., 2013), making genetic toxicology assessment an important tool in drug evaluations (Dertinger et al., 2011). Among the different mutagenic tests available, the MNT and comet assay have been widely adopted to assess the aneuploidogenic, clastogenic, and genotoxic action of drug candidates owing to their high sensitivity and low costs and execution times when compared to classical cytogenetic methods, which require metaphase observations (Dertinger et al., 2011; Frieauff et al., 2013).

The number of cells analyzed varies according to the technique employed. According to Ribeiro et al. (2003) and Frieauff et al. (2013), for the MNT in polychromatic erythrocytes, analysis of 2000 cells is recommended, whereas for the MNT in binucleated lymphocytes, 1000 cells are recommended for analysis. In the comet assay, only 100 cells need to be analyzed because of the high sensitivity of this assay in detecting free DNA fragments (Araldi et al., 2013). These values for cells analyzed are based on statistical considerations.

Micronuclei are chromosomal fragments or whole chromosomes that are lost during cell division, which can be detected as extra-chromosomal particles (Frieauff et al., 2013), indicating chromosome breakage (clastogenicity), which can lead to genomic instability, an indicator of oncogenic processes (Fenech, 2000; Araldi et al., 2013).

The evaluation of the anorectic ability of the extract of *C. ecalyculata* and its association with *S. maxima* showed a weight reduction in the male mice for the combination group Ce 500 + Sm 250 and in females for the treatment groups Ce 150, Ce 500, Ce 150 + Sm 75, and Ce 300 + Sm 150. However, the ANOVA results revealed a statistically significant anorectic effect only for the female groups of Ce 300 and Ce 500 + Sm 250, as shown in Table 4, indicating that treatment with *C. ecalyculata* alone at a dose of 300 mg/kg and in combination with *S. maxima* at the respective doses of 500 and 250 mg/kg can both contribute to higher weight reduction.

These results differ from those observed by da Silva et al. (2010), who did not observe an anorectic effect of *C. ecalyculata* at doses of 500, 1000, and 2000 mg/kg. The difference in the anorectic effect observed between the sexes is likely related to differences in the hormonal environment, which leads to alteration of gene expression and physiological responses to drugs (Guidalini and Tufik, 2007), thereby justifying the need for *in vivo* testing separately between the sexes.

The results of anorectic action in females are particularly relevant as obesity appears to be more prevalent in women (Flier and Maratos-Flier, 2013). Although obesity presents a non-Mendelian pattern of inheritance and there are at least 10 loci associated with obesity, only 3% of cases are due to a genetic component, indicating that consumption of high-energy foods and sedentary habits are the main risk factors for obesity. Therefore, the search for drugs with different targets has been actively undertaken for the treatment of appetite suppression (Kushner, 2013). The association of *C. ecalyculata* extract and *S. maximum* is an excellent candidate in this regard, because these drugs have two action pathways: 1) anorectic central action, facilitating binding to adenosine receptors, thereby promoting an extension of adrenaline (*C. ecalyculata*); and 2) peripheral action due the high protein concentration found in *S. maxima*.

The evaluation of the mutagenic potential of the treatments through the MNT, both *in vivo* and *in vitro*, and the comet assay revealed that they are safe, as they do not show clastogenicity. The tests were conducted in both mouse and human cells, as recommended by the FDA, thus providing confidence in our results. Similar results were observed by da Silva et al. (2010) at doses of 500, 1000, and 2000 mg/kg, who did not observe any genotoxic effects in cells from the peripheral blood of mice, as determined through a comet assay.

Although there are reports showing cytotoxic effects of other *Cordia* species due to the presence of pyrrolizidine alkaloids, *C. ecalyculata* does not have cytotoxic activity as it lacks this alkaloid (Rapisarda et al., 1993). Moreover, Caparroz-Assef et al. (2005) and da Silva et al. (2010) showed that *C. ecalyculata* does not cause toxicity under chronic use.

Gas chromatography, mass spectrometry, and magnetic resonance imaging analyses have shown that *C. ecalyculata* is distinguished by the presence of palmitic, myristic, stearic, oleic, linoleic acid, and high concentrations of spathulenol, a substance that inhibits the growth of *Helicobacter pylori* strain DSMZ 4867 found in human gastric samples (Menghini et al., 2008). Furthermore, *C. ecalyculata* is rich in tannins and anthocyanins, which have antioxidant properties, can modulate the expression of adipokines and prevent fat accumulation and insulin resistance type II, a problem frequently observed in obesity (Volp et al., 2008).

*S. maximum* is distinguished by the presence of alpha-tocopherol, which inhibits nuclear transcription factor kappa B (NF- $\kappa$ B). When activated by cytokines such as tumor necrosis factor- $\alpha$  and interleukin (IL)-6 and IL-8, NF- $\kappa$ B binds to the regulatory sequence of genes related to the immune processes, apoptosis, cell plasticity, and development, and is therefore related to tumorigenesis (Gilbert and Slingerland, 2013).

Cytokines secreted by preadipocytes attract monocytes and induce high concentrations of leptin, which is responsible for the conversion of monocytes to macrophages that contribute to the further production of cytokines and angiogenic factors, thereby creating a toxic microenvironment in the adipose tissue of obese individuals that is favorable for tumorigenesis (Gilbert and Slingerland, 2013). Given this scenario, the combined treatment of *C. ecalyculata* and *S. maxima* at a dose of 500 and 250 mg/kg, respectively, is recommended, as this treatment could achieve an anorectic effect while exerting antitumorigenic activity due to the presence of tannins and alpha-tocopherol.

In summary, the combined treatment of *C. ecalyculata* and *S. maxima* at a dose of 500 + 250 mg/kg proved to be a promising candidate for weight reduction therapy, and did not exhibit mutagenic action in either mouse or human cells while simultaneously conferring anti-carcinogenic action. This effect, coupled with the absence of adverse effects commonly observed in other similar treatments with drugs such as sibutramine, shows great potential of these new candidates in reducing the incidence of obesity in combination with lifestyle changes such as nutritional education and practical exercises.

## ACKNOWLEDGMENTS

The authors thank the Fundação de Amparo à Pesquisa do Estado de São Paulo (Process #2010/05649-7) for financial support and the late Dr. Tania Maria Araújo Domingues Zucchi for academic support.

## REFERENCES

- Araldi RP, Melo TC, Diniz N, Mazzuchelli-de-Souza J, et al. (2013). Bovine papillomavirus elastogenic effect analyzed in comet assay. *Biomed. Res. Int.* 2013: 630683.
- Arisawa M, Arisawa M, Fujita A, Hayashi T, et al. (1994). Cell growth inhibition of KB cells by plant extracts. *Nat. Med.* 48: 338-347.
- Batista-Filho M and Rissin A (2003). A transição nutricional no Brasil: tendências regionais e temporais. *Cad. Saúde Pública* 19: 181-191.
- Boeing H (2013). Obesity and cancer-the update 2013. *Best. Pract. Res. Clin. Endocrinol. Metab.* 27: 219-227.
- Caparroz-Assef SM, Grespan R, Batista RCF and Bersani-Amado FA (2005). Toxicity studies of *Cordia salicifolia* extract. *Acta Sci. Health Sci.* 27: 41-44.
- Carbonari K, Gonçalves L, Roth D and Moreira P (2005). Increased micronucleated cell frequency related to exposure to radiation emitted by computer cathode ray tube video display monitors. *Genet. Mol. Biol.* 28: 469-474.
- da Silva CJ, Bastos JK and Takahashi CS (2010). Evaluation of the genotoxic and cytotoxic effects of crude extracts of *Cordia ecalyculata* and *Echinodorus grandiflorus*. *J. Ethnopharmacol.* 127: 445-450.
- Delay MCEU (2002). *Arthrospira maxima* [= *Spirulina maxima* (Stiz.) Geitl., 1930] Aci Lake Strain. *J. Fish. Aquat. Sci.* 19: 241-245.
- Dertinger SD, Torous DK, Hayashi M and MacGregor JT (2011). Flow cytometric scoring of micronucleated erythrocytes: an efficient platform for assessing *in vivo* cytogenetic damage. *Mutagenesis* 26: 139-145.
- Fenech M (2000). The *in vitro* micronucleus technique. *Mutat. Res.* 455: 81-95.
- Flier JS and Maratos-Flier E (2013). Biologia da Obesidade. In: Harrison Principios Medicina Interna. 18th edn. (Longo DL, Kasper DL, Jamenson JL, Fauci AS, et al., eds.). Editora Artmed, Porto Alegre, 622-629.
- Flores M and Yamaguchi UM (2008). Teste do Micronúcleo: uma triagem para a avaliação genotóxica. *Saúde Pesq.* 1: 337-340.
- Francischi RPP, Pereira LO, Freitas CS and Klopfer M (2000). Obesidade: atualização sobre sua etiologia, morbidade e tratamento. *Rev. Nutr.* 13: 17-28.
- Frieauff W, Martus HJ, Suter W and Elhajouji A (2013). Automatic analysis of the micronucleus test in primary human lymphocytes using image analysis. *Mutagenesis* 28: 15-23.
- Gilbert CA and Slingerland JM (2013). Cytokines, obesity, and cancer: new insights on mechanisms linking obesity to cancer risk and progression. *Annu. Rev. Med.* 64: 45-57.
- Godschalk RW, Ersson C, Riso P, Porrini M, et al. (2013). DNA-repair measurements by use of the modified comet assay: an inter-laboratory comparison within the European Comet Assay Validation Group (ECVAG). *Mutat. Res.* 757: 60-67.
- Griffiths AJF, Wessler SR, Lewontin RC and Gelbart WM (2005). Um Breve Guia dos Organismos Modelo. Introdução à Genética. 8ª ed. Editora Guanabara Koogan, Rio de Janeiro, 690-691.
- Guindalini C and Tufik S (2007). Uso de microarrays na busca de perfis de expressão gênica-aplicação no estudo de fenótipos complexos. *Rev. Bras. Psiquiatr.* 29: 370-374.
- Hayashi K, Hayashi T, Morita N and Niwayama S (1990). Antiviral activity of an extract of *Cordia salicifolia* on herpes simplex virus type 1. *Planta Med.* 56: 439-443.
- Kushner RF (2013). Avaliação e Tratamento da Obesidade. In: Harrison Principios Medicina Interna. 18th edn. (Longo DL, Kasper DL, Jamenson JL, Fauci AS, et al., eds.). Editora Artmed, Porto Alegre, 629-636.
- Lehnert T, Sonntag D, Konnopka A, Riedel-Heller S, et al. (2013). Economic costs of overweight and obesity. *Best. Pract. Res. Clin. Endocrinol. Metab.* 27: 105-115.
- Matsunaga K, Sasaki S and Ohizume Y (1997). Excitatory and inhibitory effects of Paraguayan medicinal plants *Equisetum giganteum*, *Acanthospermum australe*, *Allophylus edlisand*, *Cordia salicifolia* on contraction of rabbit aorta and guinea-pig left atrium. *Nat. Med.* 51: 478-481.
- Menghini L, Epifano F, Leporini L, Pagiotti R, et al. (2008). Phytochemical investigation on leaf extract of *Cordia salicifolia* Cham. *J. Med. Food* 11: 193-194.
- Miranda MS, Cintra RG, Barros SB and Mancini FJ (1998). Antioxidant activity of the microalga *Spirulina maxima*. *Braz. J. Med. Biol. Res.* 31: 1075-1079.
- Mishima T, Murata J, Toyoshima M, Fujii H, et al. (1998). Inhibition of tumor invasion and metastasis by calcium spirulan (Ca-SP), a novel sulfated polysaccharide derived from a blue-green alga, *Spirulina platensis*. *Clin. Exp. Metastasis* 16: 541-550.
- Mittal A, Kumar PV, Banerjee S, Rao AR, et al. (1999). Modulatory potential of *Spirulina fusiformis* on carcinogen metabolizing enzymes in Swiss albino mice. *Phytother Res.* 13: 111-114.
- Ostling O and Johanson KJ (1984). Microelectrophoretic study of radiation-induced DNA damages in individual mammalian cells. *Biochem. Biophys. Res. Commun.* 123: 291-298.

- Pandey JP and Tiwari A (2010). Optimization of biomass production of *Spirulina maxima*. *J. Algal Biomass Util.* 1: 20-32.
- Ponce-Canchihumán JC, Pérez-Mendez O, Hernández-Muñoz R, Torres-Durán PV, et al. (2010). Protective effects of *Spirulina maxima* on hyperlipidemia and oxidative-stress induced by lead acetate in the liver and kidney. *Lipids Health Dis.* 9: 35.
- Rapisarda A, Ragusa S and De Pasquale A (1993). Hepatotoxic effect of the leaves of some *Cordia* species. *Acta Hort.* 332: 237-242.
- Ribeiro LR, Salvadori DMF and Marques EK (2003). Teste do Micronúcleo em Células Humanas *in vitro*. Mutagênese Ambiental. Editora Ullbra, Canoas, 355.
- Scarpato R, Verola C, Fabiani B, Bianchi V, et al. (2011). Nuclear damage in peripheral lymphocytes of obese and overweight Italian children as evaluated by the gamma-H2AX focus assay and micronucleus test. *FASEB J.* 25: 685-693.
- Schmid W (1975). The micronucleus test. *Mutat. Res.* 31: 9-15.
- Singh NP, McCoy MT, Tice RR and Schneider EL (1988). A simple technique for quantitation of low levels of DNA damage in individual cells. *Exp. Cell Res.* 175: 184-191.
- Siqueira VLD, Garcia Cortez DA, de Oliveira CE and Nakamura CV (2006). Pharmacological studies of *Cordia salicifolia* Cham in normal and diabetic rats. *Braz. Arch. Biol. Technol.* 49: 215-218.
- Tug N, Sandal S, Ozelgun B and Yilmaz B (2011). Correlation of spermogram profiles with DNA damage in sperm cells of infertile men: a comet assay study. *Gynecol. Endocrinol.* 27: 49-54.
- Volp ACP, Renhe IRT, Barra K and Stringuetta PC (2008). Flavanóides antocianinas: características e propriedades na nutrição e saúde. *Rev. Bras. Nutr. Clin.* 23: 141-149.
- Williams SC (2013). Link between obesity and cancer. *Proc. Natl. Acad. Sci. U. S. A.* 110: 8753-8754.
- Witte I, Plappert U, de WH and Hartmann A (2007). Genetic toxicity assessment: employing the best science for human safety evaluation part III: the comet assay as an alternative to *in vitro* clastogenicity tests for early drug candidate selection. *Toxicol. Sci.* 97: 21-26.
- World Health Organization (WHO) (2011). Obesity and Overweight. 2011. Obesity and Overweight - Fact Sheet #311. Available at [<http://www.who.int/mediacentre/factsheets/fs311/en/>]. Accessed 20 September 2013.